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PASSWORD:

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SESSION RESUMED IN FILE 'CPLUS' AT 16:43:39 ON 23 OCT 2007
FILE 'CPLUS' ENTERED AT 16:43:39 ON 23 OCT 2007
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COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	27.76	208.32

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-3.90	-3.90

=> file registry

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	28.23	208.79

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-3.90	-3.90

FILE 'REGISTRY' ENTERED AT 16:44:02 ON 23 OCT 2007
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STRUCTURE FILE UPDATES: 22 OCT 2007 HIGHEST RN 951207-62-8
DICTIONARY FILE UPDATES: 22 OCT 2007 HIGHEST RN 951207-62-8

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH June 29, 2007

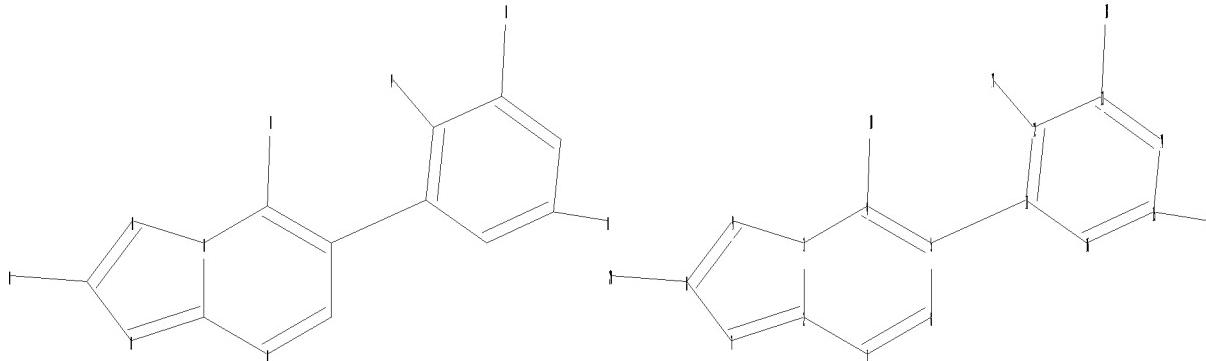
Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stndoc/properties.html>

=>

Uploading C:\Program Files\Stnexp\Queries\10 series\10594738\10594738a.str



chain nodes :
 10 17 18 19 20
 ring nodes :
 1 2 3 4 5 6 7 8 9 11 12 13 14 15 16
 chain bonds :
 4-10 5-11 8-20 12-17 13-18 15-19
 ring bonds :
 1-2 1-6 2-3 2-7 3-4 3-9 4-5 5-6 7-8 8-9 11-12 11-16 12-13 13-14 14-15
 15-16
 exact/norm bonds :
 1-2 1-6 2-3 2-7 3-4 3-9 4-5 4-10 5-6 7-8 8-9
 exact bonds :
 5-11 8-20 12-17 13-18 15-19
 normalized bonds :
 11-12 11-16 12-13 13-14 14-15 15-16

Match level :
 1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
 11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:CLASS 18:CLASS 19:CLASS
 20:CLASS

L5 STRUCTURE UPLOADED

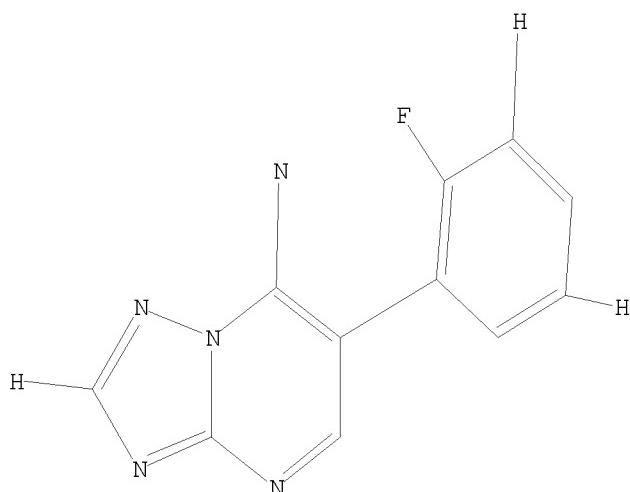
=> s 15
 SAMPLE SEARCH INITIATED 16:44:32 FILE 'REGISTRY'
 SAMPLE SCREEN SEARCH COMPLETED - 213 TO ITERATE

100.0% PROCESSED 213 ITERATIONS 46 ANSWERS
 SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
 BATCH **COMPLETE**
 PROJECTED ITERATIONS: 3385 TO 5135
 PROJECTED ANSWERS: 514 TO 1326

L6 46 SEA SSS SAM L5

=> d 15
 L5 HAS NO ANSWERS
 L5 STR



Structure attributes must be viewed using STN Express query preparation.

=> s 15 not pd>20040330

COMBINATION OF STRUCTURE AND TEXT TERMS NOT VALID

The query entered contains both search terms created by structure-building or screen commands and text search terms. L#s created via the STRUCTURE or SCREEN commands must be searched in the structures files separately from text terms or profiles. The L# answer sets from structure searches can be used in crossover searches and can be combined with text terms.

=> file caplus

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.90	209.69

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	0.00	-3.90

FILE 'CAPLUS' ENTERED AT 16:45:29 ON 23 OCT 2007

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FILE COVERS 1907 - 23 Oct 2007 VOL 147 ISS 18

FILE LAST UPDATED: 22 Oct 2007 (20071022/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply.

They are available for your review at:

<http://www.cas.org/infopolicy.html>

=> s 15 not pd>20040330
REGISTRY INITIATED
Substance data SEARCH and crossover from CAS REGISTRY in progress...
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

SAMPLE SEARCH INITIATED 16:45:34 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 213 TO ITERATE

100.0% PROCESSED 213 ITERATIONS 46 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 3385 TO 5135
PROJECTED ANSWERS: 514 TO 1326

L7 46 SEA SSS SAM L5

L8 35 L7

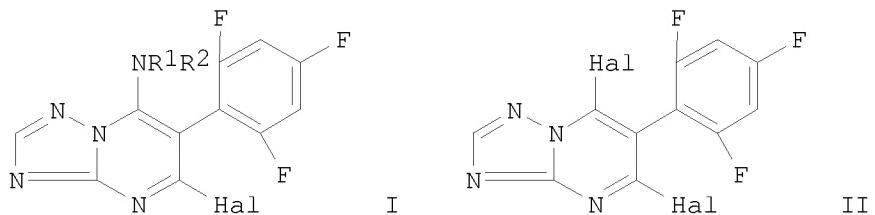
3988281 PD>20040330
(PD>20040330)
L9 3 L8 NOT PD>20040330

=> s 16 not pd>20040330
35 L6
3988281 PD>20040330
(PD>20040330)
L10 3 L6 NOT PD>20040330

=> d 110 1-3 ibib abs hitstr

L10 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 2001:719089 CAPLUS
DOCUMENT NUMBER: 135:253253
TITLE: Fungicidal trifluorophenyl-triazolopyrimidines
INVENTOR(S): Pees, Klaus-juergen; Albert, Guido
PATENT ASSIGNEE(S): American Cyanamid Co., USA
SOURCE: U.S., 11 pp.
CODEN: USXXAM
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6297251	B1	20011002	US 1999-457250	19991208
PRIORITY APPLN. INFO.:			US 1999-457250	19991208
OTHER SOURCE(S): GI	MARPAT	135:253253		



AB The compds. I [R1, R2 = H, (un)substituted alkyl, alkenyl, alkynyl, alkadienyl, haloalkyl, aryl, heteroaryl, cycloalkyl, bicycloalkyl, or heterocyclyl other than (un)substituted 2,2,2-trifluoroethyl, or R1 and R2 with interjacent N = (un)substituted heterocyclic ring; Hal = halo, provided that Hal is other than Cl when R1 = (un)branched C1-6alkyl or C3-6cycloalkyl, and R2 = H, or when R1 and R2 with interjacent N = (un)substituted piperidine] are used as active ingredients in selective fungicidal compns., which also comprise a carrier. The compds. I are prepared by treating the compds. II (Hal = halo) with an amine (R1)(R2)NH (R1, R2 as defined above).

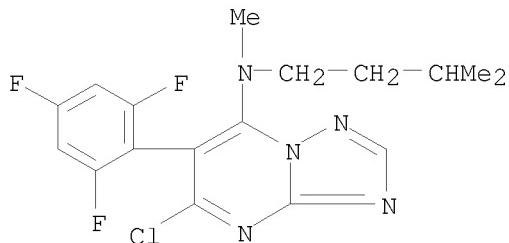
IT 214706-89-5P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)

(fungicidal trifluorophenyl-triazolopyrimidines)

RN 214706-89-5 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, 5-chloro-N-methyl-N-(3-methylbutyl)-6-(2,4,6-trifluorophenyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2001:614328 CAPLUS

DOCUMENT NUMBER: 135:176724

TITLE: Synergistic fungicidal mixtures containing azolopyrimidine and synthetic strobilurine derivatives

INVENTOR(S): Cotter, Henry Van Tuyl; May, Leslie; Reichert, Gunter; Sieverding, Ewald

PATENT ASSIGNEE(S): American Cyanamid Co., USA

SOURCE: U.S., 15 pp.

DOCUMENT TYPE: Patent

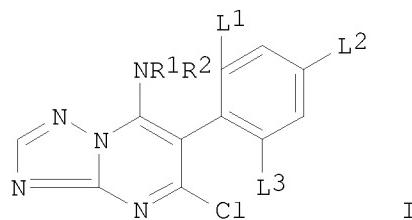
LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6277856	B1	20010821	US 1999-404910	19990924
US 6518275	B1	20030211	US 2001-809512	20010315
US 2003206968	A1	20031106	US 2002-314594	20021210
US 6699874	B2	20040302		
PRIORITY APPLN. INFO.:			US 1998-101769P	P 19980925
			US 1999-404910	A3 19990924
			US 2001-809512	A3 20010315

OTHER SOURCE(S): MARPAT 135:176724
GI



AB A synergistic fungicidal compns. comprise (a) at least one azolopyrimidine I (R1 = C1-6 alkyl, C3-6 alkenyl, C1-6 haloalkyl; or R2 = H, C1-6 alkyl; or R1R2 = C3-8 alkylene; L1 = halo; L2, L3 = H, halo) and (b) a synthetic strobilurine derivative. The compns. are used for controlling wheat leaf rust, wheat Septoria leaf blotch and/or wheat powdery mildew.

IT 355386-03-7

RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)
(synergistic fungicidal mixts. containing)

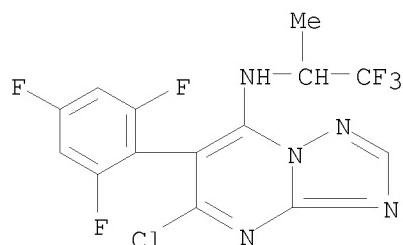
RN 355386-03-7 CAPLUS

CN Manganese, [[2-[(dithiocarboxy)amino]ethyl]carbamodithioato(2-)-
 κ S, κ S']-, mixt. with 5-chloro-N-(2,2,2-trifluoro-1-
methylethyl)-6-(2,4,6-trifluorophenyl)[1,2,4]triazolo[1,5-a]pyrimidin-7-
amine and [[2-[(dithiocarboxy)amino]ethyl]carbamodithioato(2-)-
 κ S, κ S']zinc (9CI) (CA INDEX NAME)

CM 1

CRN 214633-94-0

CMF C14 H8 Cl F6 N5

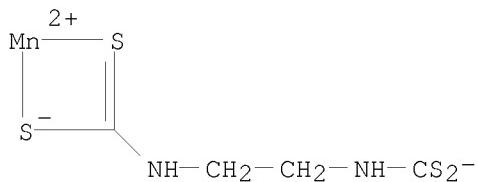


CM 2

CRN 12427-38-2

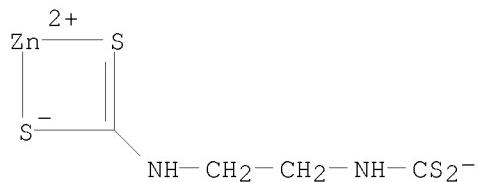
CMF C4 H6 Mn N2 S4

CCI CCS



CM 3

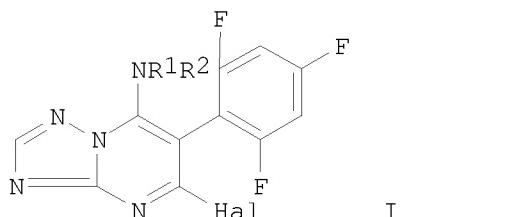
CRN 12122-67-7
 CMF C4 H6 N2 S4 Zn
 CCI CCS



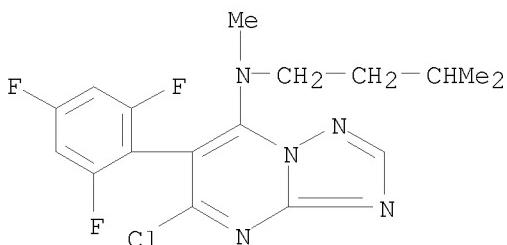
REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 2000:636212 CAPLUS
 DOCUMENT NUMBER: 133:233897
 TITLE: Preparation of fungicidal trifluorophenyl-triazolopyrimidines
 INVENTOR(S): Pees, Klaus-juergen; Albert, Guido
 PATENT ASSIGNEE(S): American Cyanamid Company, USA
 SOURCE: U.S., 10 pp.
 CODEN: USXXAM
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6117876	A	20000912	US 1998-57197	19980408
PRIORITY APPLN. INFO.:			US 1997-43816P	P 19970414
OTHER SOURCE(S): GI	MARPAT	133:233897		



AB Trifluorophenyl-triazolopyrimidine compds. I (R1 = C1-C6-alkyl or C3-C6-cycloalkyl; R2 = H; or R1 and R2 with interjacent N = piperidine, optionally substituted with one or two C1-C6-alkyls; Hal = Cl) are prepared and possess selective fungicidal activity.
 IT 214706-89-5P
 RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses (preparation as fungicide))
 RN 214706-89-5 CAPLUS
 CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, 5-chloro-N-methyl-N-(3-methylbutyl)-6-(2,4,6-trifluorophenyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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Connecting via Winsock to STN

Welcome to STN International! Enter x:X

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PASSWORD:

[USER ABORT]

Connecting via Winsock to STN

Welcome to STN International! Enter x:X

LOGINID: SSPTAJHM1624

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

* * * * * * * * * * * * * * * Welcome to STN International * * * * * * * * * * * * * * *

NEWS 1 Web Page for STN Seminar Schedule - N. America
NEWS 2 AUG 06 CAS REGISTRY enhanced with new experimental property tags
NEWS 3 AUG 06 FSTA enhanced with new thesaurus edition
NEWS 4 AUG 13 CA/CAPLus enhanced with additional kind codes for granted

patents

NEWS 5 AUG 20 CA/CAplus enhanced with CAS indexing in pre-1907 records
 NEWS 6 AUG 27 Full-text patent databases enhanced with predefined patent family display formats from INPADOCDB
 NEWS 7 AUG 27 USPATOLD now available on STN
 NEWS 8 AUG 28 CAS REGISTRY enhanced with additional experimental spectral property data
 NEWS 9 SEP 07 STN AnaVist, Version 2.0, now available with Derwent World Patents Index
 NEWS 10 SEP 13 FORIS renamed to SOFIS
 NEWS 11 SEP 13 INPADOCDB enhanced with monthly SDI frequency
 NEWS 12 SEP 17 CA/CAplus enhanced with printed CA page images from 1967-1998
 NEWS 13 SEP 17 CAplus coverage extended to include traditional medicine patents
 NEWS 14 SEP 24 EMBASE, EMBAL, and LEMBASE reloaded with enhancements
 NEWS 15 OCT 02 CA/CAplus enhanced with pre-1907 records from Chemisches Zentralblatt
 NEWS 16 OCT 19 BEILSTEIN updated with new compounds
 NEWS 17 NOV 15 Derwent Indian patent publication number format enhanced
 NEWS 18 NOV 19 WPIX enhanced with XML display format
 NEWS 19 NOV 30 ICSD reloaded with enhancements
 NEWS 20 DEC 04 LINPADOCDB now available on STN
 NEWS 21 DEC 14 BEILSTEIN pricing structure to change
 NEWS 22 DEC 17 USPATOLD added to additional database clusters
 NEWS 23 DEC 17 IMSDRUGCONF removed from database clusters and STN
 NEWS 24 DEC 17 DGENE now includes more than 10 million sequences
 NEWS 25 DEC 17 TOXCENTER enhanced with 2008 MeSH vocabulary in MEDLINE segment
 NEWS 26 DEC 17 MEDLINE and LMEDLINE updated with 2008 MeSH vocabulary
 NEWS 27 DEC 17 CA/CAplus enhanced with new custom IPC display formats
 NEWS 28 DEC 17 STN Viewer enhanced with full-text patent content from USPATOLD

NEWS EXPRESS 19 SEPTEMBER 2007: CURRENT WINDOWS VERSION IS V8.2,
 CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
 AND CURRENT DISCOVER FILE IS DATED 19 SEPTEMBER 2007.

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 NEWS IPC8 For general information regarding STN implementation of IPC 8

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* *

FILE 'HOME' ENTERED AT 18:08:30 ON 18 DEC 2007

| | | | |
|----------------------|------------|---------|--|
| => file registry | | | |
| COST IN U.S. DOLLARS | SINCE FILE | TOTAL | |
| FULL ESTIMATED COST | ENTRY | SESSION | |
| | 0.21 | 0.21 | |

FILE 'REGISTRY' ENTERED AT 18:08:45 ON 18 DEC 2007
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STRUCTURE FILE UPDATES: 17 DEC 2007 HIGHEST RN 958449-41-7
DICTIONARY FILE UPDATES: 17 DEC 2007 HIGHEST RN 958449-41-7

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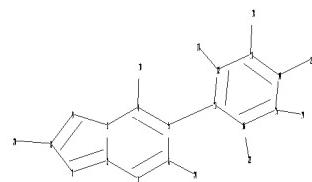
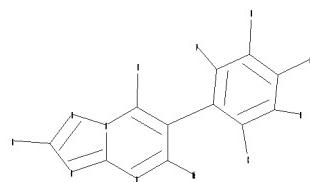
TSCA INFORMATION NOW CURRENT THROUGH June 29, 2007

Please note that search-term pricing does apply when
conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and
predicted properties as well as tags indicating availability of
experimental property data in the original document. For information
on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stndoc/properties.html>

=>
Uploading C:\Program Files\Stnexp\Queries\10 series\10594738\10594738b.str



chain nodes :

10 17 18 19 20 21 22 24

ring nodes :

1 2 3 4 5 6 7 8 9 11 12 13 14 15 16

chain bonds :

4-10 5-11 6-24 8-20 12-17 13-18 14-21 15-19 16-22

ring bonds :

1-2 1-6 2-3 2-7 3-4 3-9 4-5 5-6 7-8 8-9 11-12 11-16 12-13 13-14 14-15
15-16

exact/norm bonds :

1-2 1-6 2-3 2-7 3-4 3-9 4-5 4-10 5-6 6-24 7-8 8-9 14-21

exact bonds :

5-11 8-20 12-17 13-18 15-19 16-22

normalized bonds :

11-12 11-16 12-13 13-14 14-15 15-16

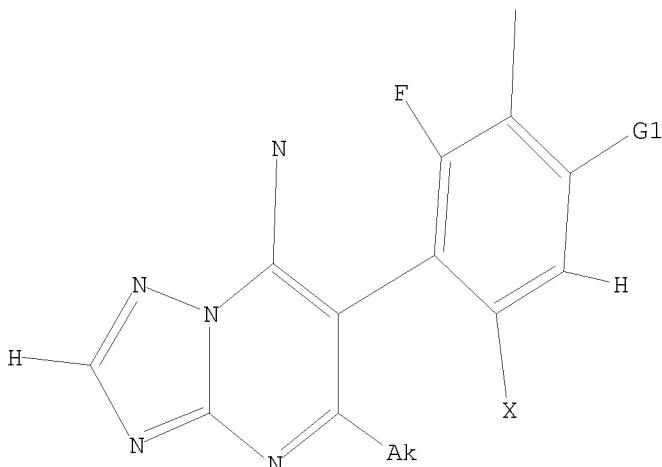
G1:H,X

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:CLASS 18:CLASS 19:CLASS
20:CLASS 21:CLASS 22:CLASS 24:CLASS

L1 STRUCTURE UPLOADED

=> d l1
L1 HAS NO ANSWERS
L1 STR



G1 H, X

Structure attributes must be viewed using STN Express query preparation.

=> s l1
SAMPLE SEARCH INITIATED 18:09:03 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 214 TO ITERATE

100.0% PROCESSED 214 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
 BATCH **COMPLETE**
PROJECTED ITERATIONS: 3403 TO 5157
PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

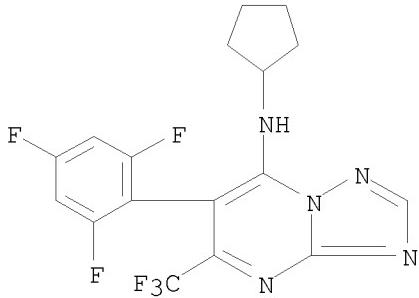
=> s l1 full
FULL SEARCH INITIATED 18:09:07 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 4191 TO ITERATE

100.0% PROCESSED 4191 ITERATIONS 17 ANSWERS
SEARCH TIME: 00.00.01

L3 17 SEA SSS FUL L1

=> d scan

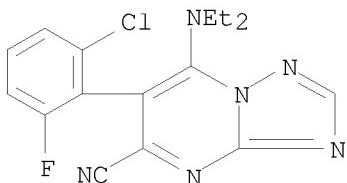
L3 17 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, N-cyclopentyl-5-(trifluoromethyl)-
6-(2,4,6-trifluorophenyl)-
MF C17 H13 F6 N5



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L3 17 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
 IN [1,2,4]Triazolo[1,5-a]pyrimidine-5-carbonitrile, 6-(2-chloro-6-fluorophenyl)-7-(diethylamino)-
 MF C16 H14 Cl F N6



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

| => file caplus | | SINCE FILE | TOTAL |
|----------------------|--|------------|---------|
| COST IN U.S. DOLLARS | | ENTRY | SESSION |
| FULL ESTIMATED COST | | 172.10 | 172.31 |

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FILE COVERS 1907 - 18 Dec 2007 VOL 147 ISS 26
FILE LAST UPDATED: 17 Dec 2007 (20071217/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply.
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<http://www.cas.org/infopolicy.html>

=> s 13
L4 9 L3

=> d 14 1-9 ibib abs hitstr

L4 ANSWER 1 OF 9 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 2006:238710 CAPLUS
DOCUMENT NUMBER: 144:292778
TITLE: Preparation of 6-phenyl-7-aminotriazolopyrimides as agrochemical fungicides
INVENTOR(S): Blettner, Carsten; Tormo, I. Blasco Jordi; Mueller, Bernd; Gewehr, Markus; Grammenos, Wassilius; Grote, Thomas; Huenger, Udo; Rheinheimer, Joachim; Schaefer, Peter; Schieweck, Frank; Schwoegler, Anja; Dietz, Jochen; Speakman, John-Bryan; Jabs, Thorsten; Strathmann, Siegfried; Schoefl, Ulrich; Scherer, Maria; Stierl, Reinhard
PATENT ASSIGNEE(S): Basf Aktiengesellschaft, Germany
SOURCE: PCT Int. Appl., 88 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

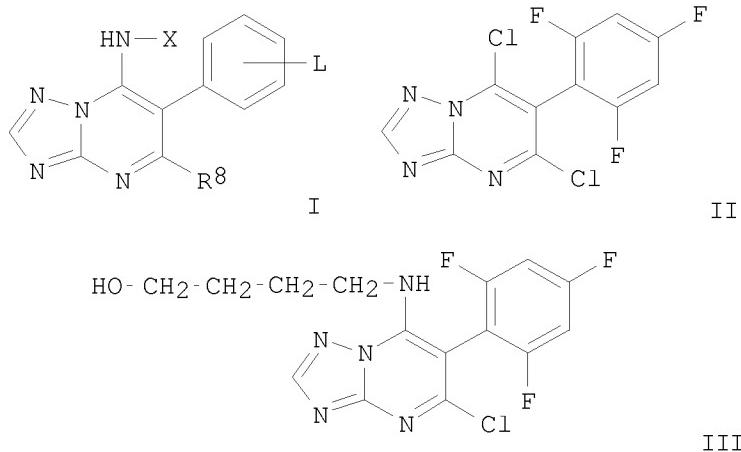
| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|-----------------------|----------|
| WO 2006027170 | A1 | 20060316 | WO 2005-EP9456 | 20050902 |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW | | | | |
| RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM | | | | |
| AU 2005281882 | A1 | 20060316 | AU 2005-281882 | 20050902 |
| CA 2577041 | A1 | 20060316 | CA 2005-2577041 | 20050902 |
| EP 1797095 | A1 | 20070620 | EP 2005-784802 | 20050902 |
| R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, HR, MK | | | | |
| CN 101014604 | A | 20070808 | CN 2005-80030105 | 20050902 |
| IN 2007KN00556 | A | 20070706 | IN 2007-KN556 | 20070214 |
| US 2007270311 | A1 | 20071122 | US 2007-661566 | 20070228 |
| KR 2007104516 | A | 20071026 | KR 2007-707910 | 20070406 |
| PRIORITY APPLN. INFO.: | | | DE 2004-102004043836A | 20040908 |

OTHER SOURCE(S):
GI

MARPAT 144:292778

WO 2005-EP9456

W 20050902



AB Title compds. I [X = CR₂R₃CR₄R₅(CR₆R₇)pYZ; R₁ = H, alkyl, haloalkyl, etc.; R₂ = alkyl, haloalkyl, cycloalkyl, etc.; R₃, R₄, R₅, R₆, R₇ = H, R₂; L = (L')m; L' = halo, alkyl, haloalkyl, etc.; R₈ = halo, CN, alkyl, etc.; Y = S, O; Z = H, alkyl, haloalkyl, etc.] were prepared. For example, condensation of 2-aminobutan-1-ol and dichloropyrimidine II afforded aminotriazolo[1,5-a]pyrimide III. In alternaria solani tomato assays, compds. I at 250 ppm, exhibited 85% protection after 5-days.

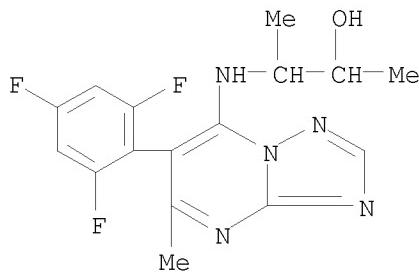
IT 879210-38-5P 879210-44-3P

RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of phenylaminotriazolo[1,5-a]pyrimides as agrochem. fungicides)

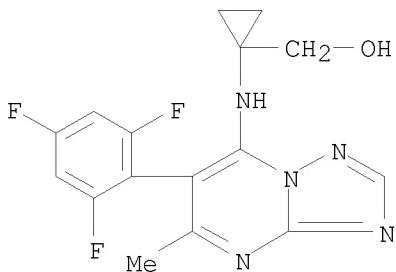
RN 879210-38-5 CAPLUS

CN 2-Butanol, 3-[[5-methyl-6-(2,4,6-trifluorophenyl)[1,2,4]triazolo[1,5-a]pyrimidin-7-yl]amino]- (CA INDEX NAME)



RN 879210-44-3 CAPLUS

CN Cyclopropanemethanol, 1-[[5-methyl-6-(2,4,6-trifluorophenyl)[1,2,4]triazolo[1,5-a]pyrimidin-7-yl]amino]- (CA INDEX NAME)

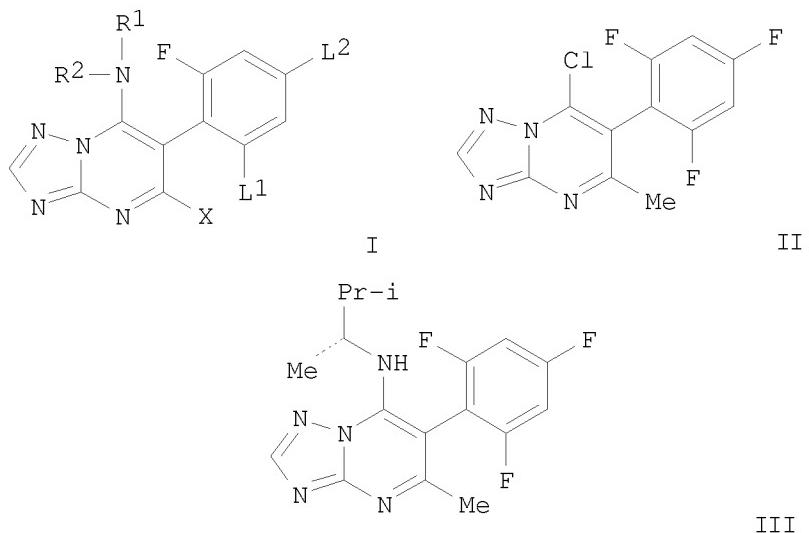


REFERENCE COUNT: 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 9 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 2005:1103781 CAPLUS
 DOCUMENT NUMBER: 143:387054
 TITLE: Preparation of 6-(2-fluorophenyl)triazolopyrimidines as agrochemical fungicides
 INVENTOR(S): Blettner, Carsten; Gewehr, Markus; Grammenos, Wassilius; Grote, Thomas; Huenger, Udo; Mueller, Bernd; Niedenbrueck, Matthias; Rheinheimer, Joachim; Schaefer, Peter; Schieweck, Frank; Schwoegler, Anja; Wagner, Oliver; Rack, Michael; Nave, Barbara; Scherer, Maria; Strathmann, Siegfried; Schoefl, Ulrich; Stierl, Reinhard
 PATENT ASSIGNEE(S): BASF Aktiengesellschaft, Germany
 SOURCE: PCT Int. Appl., 31 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|-----------------------|------------|
| WO 2005095404 | A2 | 20051013 | WO 2005-EP3208 | 20050326 |
| WO 2005095404 | A3 | 20060406 | | |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW | | | | |
| RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG | | | | |
| EP 1732927 | A2 | 20061220 | EP 2005-716387 | 20050326 |
| R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR | | | | |
| CN 1938313 | A | 20070328 | CN 2005-80010852 | 20050326 |
| BR 2005008717 | A | 20070807 | BR 2005-8717 | 20050326 |
| JP 2007530618 | T | 20071101 | JP 2007-505464 | 20050326 |
| US 2007208038 | A1 | 20070906 | US 2006-594738 | 20060929 |
| PRIORITY APPLN. INFO.: | | | DE 2004-102004016082A | 20040330 |
| | | | WO 2005-EP3208 | W 20050326 |

OTHER SOURCE(S): MARPAT 143:387054
 GI



AB Title compds. I [R1 = alkyl, haloalkyl, (un)substituted cycloalkyl, etc.; R2 = H, alkyl with provisos; L1 = Cl, F; L = H when L1 = F, F; X = alkyl] were prepared. For example, condensation of chloropyrimidine II and (2R)-3-methyl-2-butanimine afforded triazolo[1,5-a]pyrimidine III. In cucumber sphaerotheca fuliginea protection assays, 3-examples of compds. I at 250 ppm, exhibited 100% protection after 7-days.

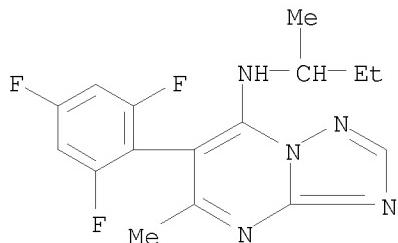
IT 866790-82-1P 866790-83-2P 866790-84-3P

RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of fluorophenyltriazolo[1,5-a]pyrimidines as agrochem. fungicides)

RN 866790-82-1 CAPLUS

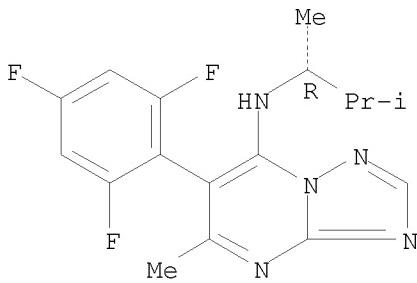
CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, 5-methyl-N-(1-methylpropyl)-6-(2,4,6-trifluorophenyl)- (CA INDEX NAME)



RN 866790-83-2 CAPLUS

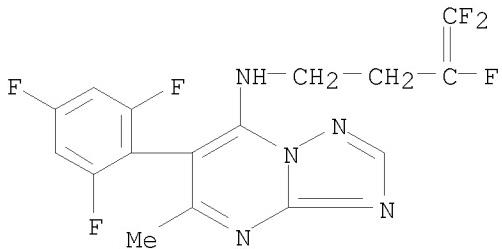
CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, N-[(1R)-1,2-dimethylpropyl]-5-methyl-6-(2,4,6-trifluorophenyl)- (CA INDEX NAME)

Absolute stereochemistry.



RN 866790-84-3 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, 5-methyl-N-(3,4,4-trifluoro-3-but enyl)-6-(2,4,6-trifluorophenyl)- (9CI) (CA INDEX NAME)



L4 ANSWER 3 OF 9 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2005:570897 CAPLUS

DOCUMENT NUMBER: 143:97391

TITLE: Preparation of 6-(2,4,6-trifluorophenyl)triazolopyrimidines for combating pathogenic fungi

INVENTOR(S): Tormo I Blasco, Jordi; Blettner, Carsten; Mueller, Bernd; Gewehr, Markus; Grammenos, Wassilius; Grote, Thomas; Rheinheimer, Joachim; Schaefer, Peter; Schieweck, Frank; Schwoegler, Anja; Wagner, Oliver; Scherer, Maria; Strathmann, Siegfried; Schoefl, Ulrich; Stierl, Reinhard

PATENT ASSIGNEE(S): BASF Aktiengesellschaft, Germany

SOURCE: PCT Int. Appl., 34 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: German

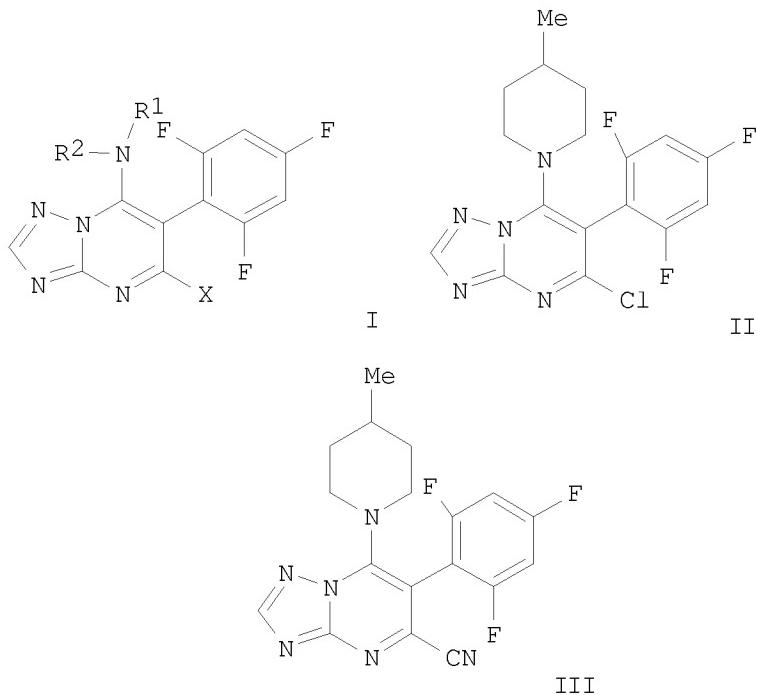
FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|-----------------|----------|
| WO 2005058900 | A1 | 20050630 | WO 2004-EP13063 | 20041118 |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW | | | | |
| RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, | | | | |

NE, SN, TD, TG
 EP 1689750 A1 20060816 EP 2004-797968 20041118
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
 IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK, IS
 CN 1886404 A 20061227 CN 2004-80034578 20041118
 BR 2004016849 A 20070227 BR 2004-16849 20041118
 JP 2007512276 T 20070517 JP 2006-540311 20041118
 US 2007149515 A1 20070628 US 2006-579144 20060515
 PRIORITY APPLN. INFO.: DE 2003-10355387 A 20031125
 WO 2004-EP13063 W 20041118

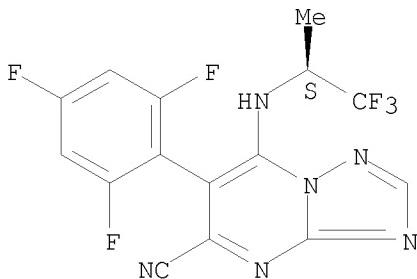
OTHER SOURCE(S): MARPAT 143:97391
 GI



AB Title compds. I [R1 = alkyl, haloalkyl, cycloalkyl, etc.; R2 = H, or together with R1 with provisos; X = CN, alkoxy, alkenyloxy, etc.] were prepared. For example, tetrabutylammonium cyanide mediated nitrilation of chloropyrimidine II afforded triazolo[1,5-a]pyrimidine III. In *sphaerotheca fuliginea* protection assays, 4-examples of compds. I, at 63 ppm application, after 7-days exhibited 100% protection.

IT 856543-22-1P
 RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (preparation of trifluorophenyltriazolo[1,5-a]pyrimidines for combating pathogenic fungi)
 RN 856543-22-1 CAPLUS
 CN [1,2,4]Triazolo[1,5-a]pyrimidine-5-carbonitrile, 7-[[[(1S)-2,2,2-trifluoro-1-methylethyl]amino]-6-(2,4,6-trifluorophenyl)- (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 4 OF 9 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2004:857602 CAPLUS

DOCUMENT NUMBER: 141:332222

TITLE: Methods for the production and use of

7-(alkynylamino)triazolo[4,5-d]pyrimidines and agents

containing them useful for combating harmful fungi

Tormo I Blasco, Jordi; Blettner, Carsten; Mueller, Bernd; Gewehr, Markus; Grammenos, Wassilius; Grote, Thomas; Gypser, Andreas; Rheinheimer, Joachim; Schaefer, Peter; Schieweck, Frank; Schwoegler, Anja; Scherer, Maria; Strathmann, Siegfried; Schoefl, Ulrich; Stierl, Reinhard

PATENT ASSIGNEE(S): BASF Aktiengesellschaft, Germany

SOURCE: PCT Int. Appl., 36 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: German

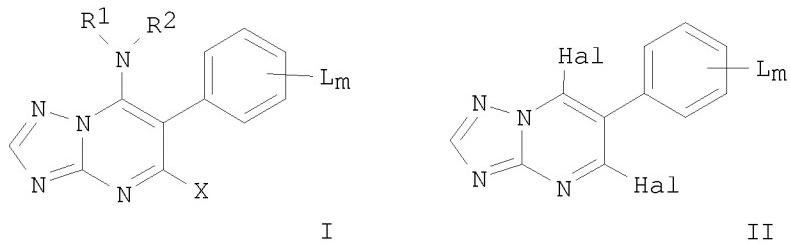
FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|--|------|----------|------------------|------------|
| WO 2004087706 | A1 | 20041014 | WO 2004-EP3346 | 20040330 |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE,
ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI,
SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN,
TD, TG | | | | |
| AU 2004226253 | A1 | 20041014 | AU 2004-226253 | 20040330 |
| CA 2520718 | A1 | 20041014 | CA 2004-2520718 | 20040330 |
| EP 1613633 | A1 | 20060111 | EP 2004-724256 | 20040330 |
| R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK | | | | |
| BR 2004008864 | A | 20060411 | BR 2004-8864 | 20040330 |
| CN 1768062 | A | 20060503 | CN 2004-80009242 | 20040330 |
| JP 2006522046 | T | 20060928 | JP 2006-504913 | 20040330 |
| US 2006211711 | A1 | 20060921 | US 2005-550571 | 20050923 |
| IN 2005CN02849 | A | 20070720 | IN 2005-CN2849 | 20051102 |
| PRIORITY APPLN. INFO.: | | | DE 2003-10314930 | A 20030402 |
| | | | WO 2004-EP3346 | W 20040330 |

OTHER SOURCE(S):
GI

CASREACT 141:332222; MARPAT 141:332222



AB 7-(Alkynylamino)triazolopyrimidines I [L = halogen, C1-6-alkyl, C1-6-halogenalkyl, C1-6-alkoxy, NH₂, NHR, NR₂, cyano, S(O)nA1 or C(O)A2; R = C1-8-alkyl, C1-8-alkylcarbonyl; A1 = hydrogen, hydroxy, C1-8-alkyl, C1-8-alkylamino, di(C1-8-alkyl)amino; n = 0, 1 or 2; A2 = C2-8-alkenyl, C1-8-alkoxy, C1-6-halogenalkoxy or A1; m = 1, 2, 3, 4 or 5 (whereby at least one group L is present in an ortho-position to the bond with the triazolopyrimidine skeleton); X = halogen, cyano, C1-4-alkyl, C1-4-haloalkyl, C1-4-alkoxy; R1 = hydrogen, C1-4-alkyl; R2 = (un)substituted C3-10-alkynyl]. The invention also relates to methods for the production of said compds., agents containing said compds. and the use thereof

to combat harmful phytopathogenic fungi. The procedure for the preparation of I is characterized by: reaction of halotriazolopyrimidines II (Hal = halogen) with R₁R₂NH. Thus, triazolopyrimidine I [R₁ = H, R₂ = CH₂C.tpbond.CH, X = Cl, L₃ = F3-2,4,6] was prepared from 5,7-Dichloro-6-(2,4,6-trifluorophenyl)[1,2,4]triazolo[1,5-a]pyrimidine (II;) via amination with HC.tpbond.CCH₂NH₂ in CH₂C₁₂ containing Et₃N. The inhibitory activity of I were determined [after 5 d I (R₁ = H, R₂ = CH₂C.tpbond.CCH₂Cl, X = Cl, L₃ = F3-2,4,6; R₁ = H, R₂ = CMe₂C.tpbond.CH, X = Cl, L₃ = F3-2,4,6) had decreased the activity of *Alternaria solani* (Tomato dry spot disease) and *Puccinia recondita* (wheat brown rust) to 3%].

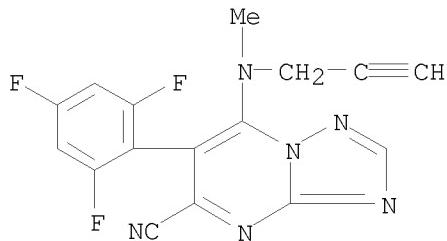
IT 773879-70-2P 773879-72-4P

RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of (alkynylamino)triazolopyrimidines for use in combating harmful phytopathogenic fungi)

RN 773879-70-2 CAPLUS

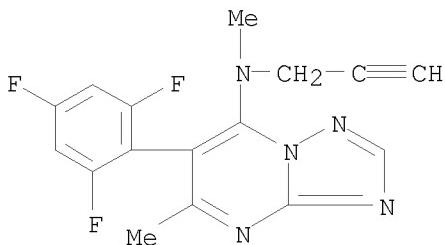
CN [1,2,4]Triazolo[1,5-a]pyrimidine-5-carbonitrile, 7-(methyl-2-propynylamino)-6-(2,4,6-trifluorophenyl)- (9CI) (CA INDEX NAME)



RN 773879-72-4 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, N,5-dimethyl-N-2-propynyl-6-

(2,4,6-trifluorophenyl)- (9CI) (CA INDEX NAME)

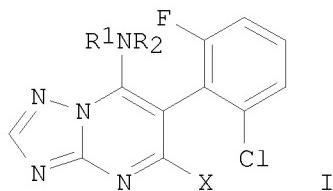


REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 5 OF 9 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 2002:814135 CAPLUS
DOCUMENT NUMBER: 137:325429
TITLE: Preparation of 6-(2-chloro-6-fluorophenyl)-triazolopyrimidines as agrochemical fungicides
INVENTOR(S): Tormo i Blasco, Jordi; Sauter, Hubert; Mueller, Bernd; Gewehr, Markus; Grammenos, Wassilius; Grote, Thomas; Gypser, Andreas; Rheinheimer, Joachim; Rose, Ingo; Schaefer, Peter; Schiweck, Frank; Ammermann, Eberhard; Strathmann, Siegfried; Lorenz, Gisela; Stierl, Reinhard
PATENT ASSIGNEE(S): Basf Aktiengesellschaft, Germany
SOURCE: PCT Int. Appl., 32 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|-----------------|------------|
| WO 2002083677 | A1 | 20021024 | WO 2002-EP3830 | 20020406 |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW | | | | |
| RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG | | | | |
| AU 2002257758 | A1 | 20021028 | AU 2002-257758 | 20020406 |
| EP 1381610 | A1 | 20040121 | EP 2002-727534 | 20020406 |
| EP 1381610 | B1 | 20040825 | | |
| R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR | | | | |
| JP 2004526767 | T | 20040902 | JP 2002-581432 | 20020406 |
| AT 274518 | T | 20040915 | AT 2002-727534 | 20020406 |
| ES 2225784 | T3 | 20050316 | ES 2002-2727534 | 20020406 |
| US 2004110751 | A1 | 20040610 | US 2003-474461 | 20031008 |
| US 7071334 | B2 | 20060704 | | |
| PRIORITY APPLN. INFO.: | | | EP 2001-109010 | A 20010411 |
| | | | WO 2002-EP3830 | W 20020406 |

OTHER SOURCE(S): MARPAT 137:325429
GI



AB The title compds. [I; R1, R2 = H, alkyl, (un)substituted Ph, heterocyclyl, etc.; or NR1R2 = (un)substituted 5-6 membered heterocyclic ring; X = CN, alkoxy, haloalkoxy, alkenyloxy], useful for combating phytopathogenic fungi, were prepared Thus, treating I [NR1R2 = 4-methylpiperidino; X = Cl] with NaOMe in MeOH afforded I [NR1R2 = 4-methylpiperidino; X = OMe]. The tomato plants (infested by Alternaria solani) which had been treated with 63 ppm of the latter showed an infection of up 3%, whereas the untreated plants were infected to 100%.

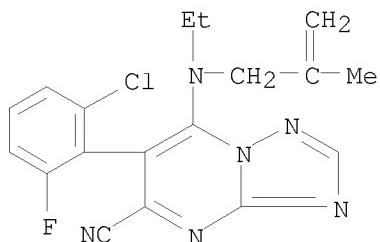
IT 473465-98-4P 473465-99-5P 473466-01-2P

RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of 6-(2-chloro-6-fluorophenyl)triazolopyrimidines as agrochem. fungicides)

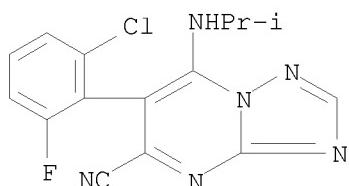
RN 473465-98-4 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidine-5-carbonitrile, 6-(2-chloro-6-fluorophenyl)-7-[ethyl(2-methyl-2-propenyl)amino]- (9CI) (CA INDEX NAME)



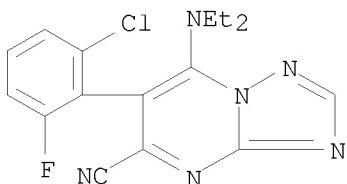
RN 473465-99-5 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidine-5-carbonitrile, 6-(2-chloro-6-fluorophenyl)-7-[(1-methylethyl)amino]- (CA INDEX NAME)



RN 473466-01-2 CAPLUS

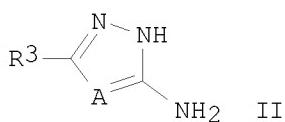
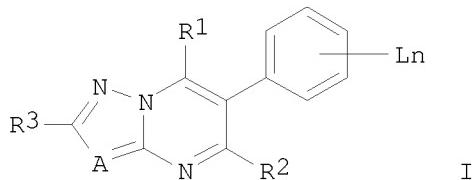
CN [1,2,4]Triazolo[1,5-a]pyrimidine-5-carbonitrile, 6-(2-chloro-6-fluorophenyl)-7-(diethylamino)- (CA INDEX NAME)



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 6 OF 9 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 2002:807309 CAPLUS
 DOCUMENT NUMBER: 137:325424
 TITLE: Preparation of 5-(haloalkyl)azolopyrimidines and their use as pesticides
 INVENTOR(S): Miyahara, Osamu; Hamamura, Hiroshi; Hirai, Yukio; Yokota, Yori
 PATENT ASSIGNEE(S): Nippon Soda Co., Ltd., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 35 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------------------------|------------|-------------------|---|-------------------------------|
| -----
JP 2002308879 | -----
A | -----
20021023 | -----
JP 2001-115989
JP 2001-115989 | -----
20010413
20010413 |
| PRIORITY APPLN. INFO.: | | | | |
| OTHER SOURCE(S): | MARPAT | 137:325424 | | |
| GI | | | | |



AB Title compds. I [R1 = H, OH, halo, C1-8 (halo)alkyl, C2-8 alkenyl, C2-8 alkynyl, C3-8 cycloalkyl, (un)substituted heterocyclyl, (un)substituted aryl, amino, etc.; R2 = C1-8 haloalkyl; R3 = H, C1-4 alkyl,

(un)substituted aryl; L = halo, C1-4 alkyl, C1-3 haloalkyl, C1-4 alkoxy, C1-3 haloalkoxy; n = 0-5; A = N, CH] or their salts are useful as marine antifouling agents, insecticides, acaricides (no data), and agrochem fungicides. I (R1 = OH; R2, R3, L, n, A = same as above) are prepared by treatment of R2COCH(C6H5-nLn)CO2R4 [R2, L, n = same as above; R4 = C1-4 alkyl, (un)substituted Ph] with azoles II (R3, A = same as above). Thus, I (R1 = OH, R2 = CF₃, R3 = H, Ln = 2-Cl-6-F-C₆H₃, A = N) was chlorinated with POC₁₃ to give the corresponding chloride with 52% yield, which was condensed with 4-piperidine to afford 85% I (R1 = 4-piperidino, R2 = CF₃, R3 = H, Ln = 2-Cl-6-F-C₆H₃, A = N). The product showed ≥75% antifungal activity against *Venturia inaequalis*.

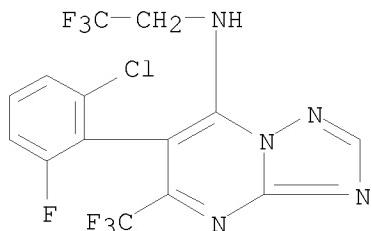
IT 473435-13-1P 473435-15-3P 473435-26-6P
473435-28-8P

RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of 5-(haloalkyl)azolopyrimidines as pesticides)

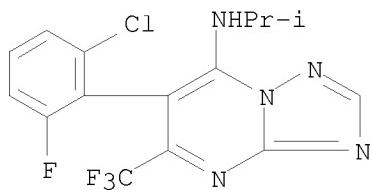
RN 473435-13-1 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, 6-(2-chloro-6-fluorophenyl)-N-(2,2,2-trifluoroethyl)-5-(trifluoromethyl)- (CA INDEX NAME)



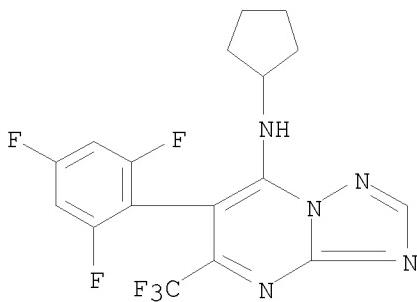
RN 473435-15-3 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, 6-(2-chloro-6-fluorophenyl)-N-(1-methylethyl)-5-(trifluoromethyl)- (CA INDEX NAME)



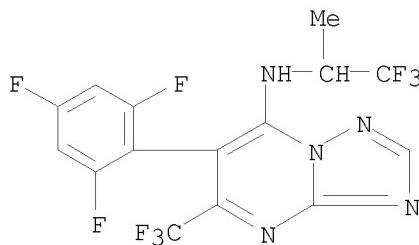
RN 473435-26-6 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, N-cyclopentyl-5-(trifluoromethyl)-6-(2,4,6-trifluorophenyl)- (CA INDEX NAME)



RN 473435-28-8 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, 5-(trifluoromethyl)-N-(2,2,2-trifluoro-1-methylethyl)-6-(2,4,6-trifluorophenyl)- (CA INDEX NAME)



L4 ANSWER 7 OF 9 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2002:31452 CAPLUS

DOCUMENT NUMBER: 136:96032

TITLE: Substituted triazolopyrimidines as anticancer agents

INVENTOR(S): Schmitt, Mark R.; Kirsch, Donald R.; Harris, Jane E.; Beyer, Carl F.; Pees, Klaus-Juergen; Carter, Paul; Pfrengle, Waldemar; Albert, Guido

PATENT ASSIGNEE(S): American Home Products Corporation, USA

SOURCE: PCT Int. Appl., 405 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|-----------------|----------|
| WO 2002002563 | A2 | 20020110 | WO 2001-US20672 | 20010628 |
| WO 2002002563 | A3 | 20030103 | | |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW | | | | |
| RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG | | | | |
| CA 2413802 | A1 | 20020110 | CA 2001-2413802 | 20010628 |
| BR 2001012038 | A | 20030401 | BR 2001-12038 | 20010628 |
| EP 1307200 | A2 | 20030507 | EP 2001-952295 | 20010628 |

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|--|----|----------|-----------------|-------------|
| R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
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| HU 2003000798 | A2 | 20030728 | HU 2003-798 | 20010628 |
| JP 2004502691 | T | 20040129 | JP 2002-507815 | 20010628 |
| NZ 523807 | A | 20040924 | NZ 2001-523807 | 20010628 |
| CN 1592624 | A | 20050309 | CN 2001-812055 | 20010628 |
| US 2002068744 | A1 | 20020606 | US 2001-895975 | 20010629 |
| BG 107277 | A | 20040130 | BG 2002-107277 | 20021115 |
| MX 2002PA11913 | A | 20030422 | MX 2002-PA11913 | 20021202 |
| NO 2002006195 | A | 20030227 | NO 2002-6195 | 20021223 |
| IN 2003KN00001 | A | 20050311 | IN 2003-KN1 | 20030101 |
| ZA 2003000793 | A | 20040720 | ZA 2003-793 | 20030129 |
| IN 2007KN00659 | A | 20070706 | IN 2007-KN659 | 20070222 |
| PRIORITY APPLN. INFO.: | | | US 2000-215585P | P 20000630 |
| | | | WO 2001-US20672 | W 20010628 |
| | | | IN 2003-KN1 | A3 20030101 |

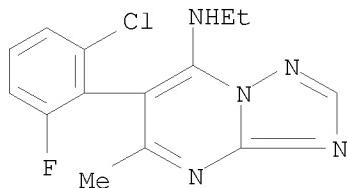
OTHER SOURCE(S): MARPAT 136:96032

AB A method is provided for treating or inhibiting the growth of cancerous tumor cells and associated diseases in a mammal in need thereof which comprises administering to the mammal an effective amount of a substituted triazolopyrimidine derivative or a pharmaceutically acceptable salt thereof. Also provided is a method for treating or inhibiting the growth of cancerous tumor cells and associated diseases in a mammal in need thereof by interacting with tubulin and microtubules and promoting microtubule polymerization which comprises administering to the mammal an effective amount of a substituted triazolopyrimidine derivative or a pharmaceutically acceptable salt thereof.

IT 220482-12-2
 RL: DMA (Drug mechanism of action); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (triazolopyrimidine derivs. as anticancer agents)

RN 220482-12-2 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, 6-(2-chloro-6-fluorophenyl)-N-ethyl-5-methyl- (CA INDEX NAME)



L4 ANSWER 8 OF 9 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1999:761522 CAPLUS
 DOCUMENT NUMBER: 131:351347
 TITLE: Preparation of fungicidal 5-alkyl-triazolopyrimidines
 INVENTOR(S): Pfrengle, Waldemar
 PATENT ASSIGNEE(S): American Cyanamid Company, USA
 SOURCE: U.S., 9 pp.
 CODEN: USXXAM
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

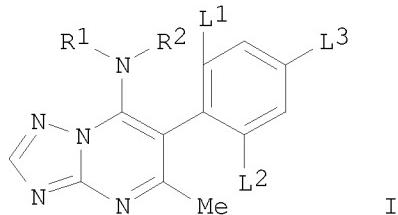
| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------------|------|-------|-----------------|-------|
| ----- | ---- | ----- | ----- | ----- |

US 5994360
PRIORITY APPLN. INFO.:
OTHER SOURCE(S):
GI

A 19991130
MARPAT 131:351347

US 1998-115496
US 1997-52407P

P 19980714
P 19970714



I

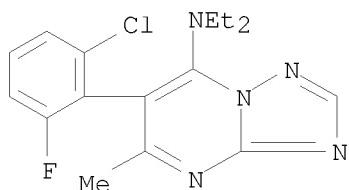
AB The title compds. [I; NR1R2 = piperidino, 4-methylpiperidino; L1-L3 = H, F, Cl (at least one of which being F or Cl) which show selective fungicidal activity, were prepared. Thus, reacting 6-(2-chloro-6-fluorophenyl)-5-chloro-7-(4-methylpiperidin-1-yl)-[1,2,4]triazolo[1,5-a]pyrimidine with di-Et malonate in the presence of NaH in MeCN followed by treatment of the resulting di-Et [6-(2-chloro-6-fluorophenyl)-7-(4-methylpiperidin-1-yl)-[1,2,4]triazolo[1,5-a]pyrimidin-yl]malonate with concentrate HCl afforded I [R1R2 = (CH₂)₂CH(Me)(CH₂)₂; L1 = Cl; L2 = F; L3 = H] which showed ED₅₀ > 90 at 0.2 mg/mL in test with Alternaria solani.

IT 220482-11-1P 220482-12-2P

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of fungicidal 5-alkyl-triazolopyrimidines)

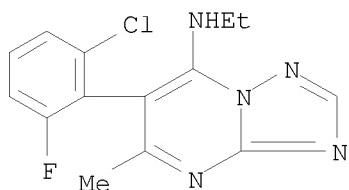
RN 220482-11-1 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, 6-(2-chloro-6-fluorophenyl)-N,N-diethyl-5-methyl- (CA INDEX NAME)



RN 220482-12-2 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, 6-(2-chloro-6-fluorophenyl)-N-ethyl-5-methyl- (CA INDEX NAME)



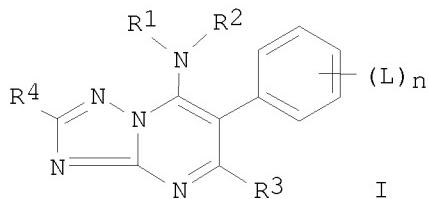
REFERENCE COUNT:

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THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 9 OF 9 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1999:106975 CAPLUS
 DOCUMENT NUMBER: 130:168390
 TITLE: Preparation of 5-alkyltriazolopyrimidines, and
 agrochemical bactericidal and fungicidal compositions
 containing them
 INVENTOR(S): Pfrengle, Waldermar Franz Augustin
 PATENT ASSIGNEE(S): American Cyanamid Co., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 13 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

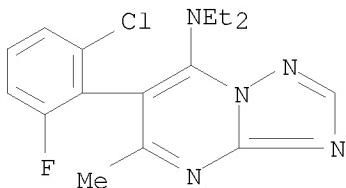
| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------------------------|--------|------------|-----------------|------------|
| JP 11035581 | A | 19990209 | JP 1998-208531 | 19980709 |
| FR 2765875 | A1 | 19990115 | FR 1998-8423 | 19980701 |
| FR 2765875 | B1 | 19991119 | | |
| PRIORITY APPLN. INFO.: | | | US 1997-892495 | A 19970714 |
| OTHER SOURCE(S): | MARPAT | 130:168390 | | |
| GI | | | | |



AB The title compds. I [R1 = (un)substituted alkyl, alkenyl, alkynyl, aryl, heteroaryl, etc.; R2 = H, (un)substituted alkyl, alkenyl, alkynyl, aryl, heteroaryl, etc.; R1NR2 may form (un)substituted heterocycll; R3 = alkyl, R4 = H, alkyl, aryl; L = halo, (un)substituted alkyl, alkoxy; A = N, CR5; R5 = similar group as shown in R4; n = 0-5] are claimed. I (R1, R2, R4, A, L, n = same as above; R3 = Me) are prepared by treatment of 5-haloazopyrimidines I (R1, R2, R4, A, L, n = same as above; R3 = halo) with alkyl malonate in the presence of bases, then heating the resulting modified malonate esters with acids. I [R1NR2 = 4-methylpiperidin-1-yl, R3 = CH(CO₂Et)₂, R4 = H, A = N, Ln = 2-Cl, 6-F] (0.5 g) was treated with concentrated HCl at 80° for 24 h to give 0.27 g I (R1NR2, R4, A, Ln = same as above, R3 = Me), which showed strong antimicrobial activities.

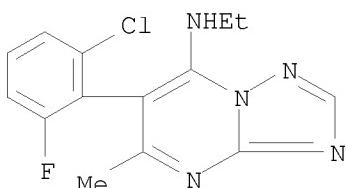
IT 220482-11-1P 220482-12-2P
 RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); IMF (Industrial manufacture); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of 5-alkyltriazolopyrimidines as agrochem. bactericides and fungicides)

RN 220482-11-1 CAPLUS
 CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, 6-(2-chloro-6-fluorophenyl)-N,N-diethyl-5-methyl- (CA INDEX NAME)



RN 220482-12-2 CAPLUS

CN [1,2,4]Triazolo[1,5-a]pyrimidin-7-amine, 6-(2-chloro-6-fluorophenyl)-N-ethyl-5-methyl- (CA INDEX NAME)



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| => file registry | | | |
| COST IN U.S. DOLLARS | SINCE FILE | TOTAL | |
| | ENTRY | SESSION | |
| FULL ESTIMATED COST | 55.89 | 228.20 | |
| DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) | SINCE FILE | TOTAL | |
| | ENTRY | SESSION | |
| CA SUBSCRIBER PRICE | -7.02 | -7.02 | |

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 DICTIONARY FILE UPDATES: 17 DEC 2007 HIGHEST RN 958449-41-7

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<http://www.cas.org/support/stngen/stndoc/properties.html>

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normalized bonds :
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G1:H,X

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Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:CLASS 18:CLASS 19:CLASS
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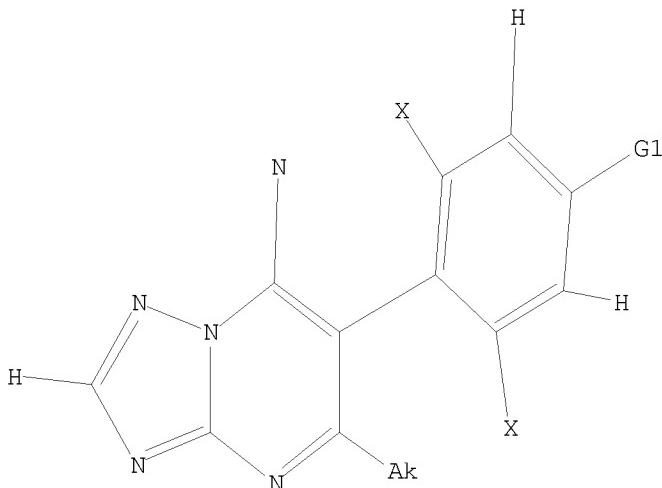
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G1 H,X

Structure attributes must be viewed using STN Express query preparation.

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100.0% PROCESSED 245 ITERATIONS
SEARCH TIME: 00.00.01

0 ANSWERS

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**

PROJECTED ITERATIONS: 3961 TO 5839

PROJECTED ANSWERS: 0 TO 0

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=> s 15 full

FULL SEARCH INITIATED 18:20:21 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 4976 TO ITERATE

100.0% PROCESSED 4976 ITERATIONS
SEARCH TIME: 00.00.01

17 ANSWERS

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COST IN U.S. DOLLARS

FULL ESTIMATED COST

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| DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) | SINCE FILE ENTRY | TOTAL SESSION |
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| CA SUBSCRIBER PRICE | 0.00 | -7.02 |

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 FILE LAST UPDATED: 17 Dec 2007 (20071217/ED)

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 L3 17 S L1 FULL

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 L7 17 S L5 FULL

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